

AMENDMENTS TO THE SPECIFICATION

Please amend the specification as follows:

1. Paragraph on page 5 starting at line 4 and ending at line 19:

wherein R_1 and R_2 , and R_3 and R_4 taken together each independently comprise a substituted or unsubstituted cyclic or polycyclic aryl or heteroaromatic moiety; wherein m is 1, 2, or 3; wherein n is 0 or 1; wherein Z or Y each independently comprise CR_{2-} , -C(R)_{2-} , wherein each occurrence of the functional moiety R , is independently selected from the group consisting of hydrogen and methyl; NR , wherein R is selected from the group consisting of hydrogen and methyl; O ; S ; or Se ; wherein X is a non-coordinating negative counter ion including, but not limited to BF_4 , PF_6 , ClO_4 , TsO , I , Br ; and wherein R_5 or R_6 each independently comprise lower alkyl, a chiral reagent (CR) or a chiral reagent and linker (L-CR), whereby said chiral reagent is attached to the detecting agent via the linker, with the proviso that at least one of R_5 or R_6 is a chiral reagent (CR) or a chiral reagent and linker (L-CR); and

wherein the chiral reagent (CR) for the first chiral detecting reagent (CDR) and for the second chiral detecting reagent (CDR) in a pair are enantiomers, and wherein each of said chiral detecting reagents in a set is capable of selectively reacting with one enantiomeric reaction product over the other enantiomeric reaction product in a sample of reaction products and is capable of being uniquely identified.

2. Paragraph on page 6 starting at line 4 and ending at line 14:

wherein R_1 and R_2 , and R_3 and R_4 taken together each independently comprise a substituted or unsubstituted cyclic or polycyclic aryl or heteroaromatic moiety; wherein m is 1, 2, or 3; wherein n is 0 or 1; wherein Z or Y each independently comprise CR_{2-} , -C(R)_{2-} , wherein each occurrence of the functional moiety R , is independently selected from the group consisting of hydrogen and methyl; NR , wherein R is selected from the group consisting of hydrogen and methyl; O ; S ; or Se ; wherein X is a non-coordinating negative counter ion including, but not limited to BF_4 , PF_6 , ClO_4 , TsO , I , Br ; and wherein R_5 or R_6 each independently comprise lower alkyl, a chiral reagent (CR) or a chiral reagent and linker (L-CR), whereby said chiral reagent is attached to the detecting agent via the linker, with the proviso that at least one of R_5 or R_6 is a chiral reagent (CR) or a chiral reagent and linker (L-CR).

3. Paragraph on page 7 starting at line 5 and ending at line 20:

wherein R_1 and R_2 , and R_3 and R_4 taken together each independently comprise a substituted or unsubstituted cyclic or polycyclic aryl or heteroaromatic moiety; wherein m is 1, 2, or 3; wherein n is 0 or 1; wherein Z or Y each independently comprise CR_{2n} , $-\text{C(R)}_{2n}$, wherein each occurrence of the functional moiety R , is independently selected from the group consisting of hydrogen and methyl; NR , wherein R is selected from the group consisting of hydrogen and methyl; O ; S ; or Se ; wherein X is a non-coordinating negative counter ion including, but not limited to BF_4 , PF_6 , ClO_4 , TsO , I , Br ; and wherein R_5 or R_6 each independently comprise lower alkyl, a chiral reagent (CR) or a chiral reagent and linker (L-CR), whereby said chiral reagent is attached to the detecting agent via the linker, with the proviso that at least one of R_5 or R_6 is a chiral reagent (CR) or a chiral reagent and linker (L-CR); and

wherein the chiral reagent (CR) for the first chiral detecting reagent (CDR) and for the second chiral detecting reagent (CDR) in a pair are enantiomers, and wherein each of said chiral detecting reagents is capable of selectively reacting with one enantiomeric reaction product over the other enantiomeric reaction product in a pair and is capable of being uniquely identified.

4. Paragraph on page 23 starting at line 6 and ending at line 23:

wherein R_1 and R_2 , and R_3 and R_4 taken together each independently comprise a substituted or unsubstituted cyclic or polycyclic aryl or heteroaromatic moiety; wherein m is 1, 2, or 3; wherein n is 0 or 1; wherein Z or Y each independently comprise CR_{2n} , $-\text{C(R)}_{2n}$, wherein each occurrence of the functional moiety R is independently selected from the group consisting of hydrogen and methyl; NR , wherein R is selected from the group consisting of hydrogen and methyl; O ; S ; or Se ; wherein X is a non-coordinating negative counter ion including, but not limited to BF_4 , PF_6 , ClO_4 , TsO , I , Br ; and wherein R_5 or R_6 each independently comprise lower alkyl, a chiral reagent (CR) or a chiral reagent and linker (L-CR), whereby said chiral reagent is attached to the detecting agent via the linker. In certain embodiments, the linker moiety comprises $-(\text{CH})_p-(\text{CO})-$, wherein p is 1-5, most preferably 4, and the chiral agent comprises a chiral acylating agent, as described in certain embodiments herein. In certain embodiments, the chiral agent comprises a chiral acylating agent having the general structure: $-(\text{NH})-(\text{CHR}_x)-\text{COOH}$, where R_x comprises a chiral amino acid residue. In certain other embodiments of the present invention, the chiral reagent (CR) or the chiral reagent and linker (L-CR) can be attached

at any one of R_1 - R_6 , or as substitutions of other moieties thereof attached at R_1 - R_6 (e.g, the chiral reagent or chiral reagent and linker can be attached via an aryl moiety or other functional ~~group~~ group).